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| <p>(54) Title: <b>BIODEGRADABLE POLYMERIC ENDOLUMINAL SEALING</b></p> <div data-bbox="435 1272 1317 1507" data-label="Image"> </div> <p>(57) Abstract</p> <p>A novel process for paving or sealing the interior surface of a tissue lumen by entering the interior of the tissue lumen and applying a polymer to the interior surface of the tissue lumen. This is accomplished using a catheter which delivers the polymer to the tissue lumen (134) and causes it to conform to the interior surface of lumen. The polymer can be delivered to the lumen as a monomer or prepolymer solution, or as an at least partially preformed layer on an expansible member (133).</p> |                  |                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |

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Description--Biodegradable Polymeric Endoluminal Sealing--Background of the Invention

5 This invention relates to a novel method for the  
in vivo paving and sealing of the interior of organs  
or organ components and other tissue cavities, and to  
apparatus and partially pre-formed polymeric products  
for use in this method. The tissues involved may be  
10 those organs or structures having hollow or tubular  
geometry, for example blood vessels such as arteries  
or veins, in which case the polymeric products are  
deposited within the naturally occurring lumen.  
Alternatively, the tissue may be a normally solid  
15 organ in which a cavity has been created either as a  
result of an intentional surgical procedure or an  
accidental trauma. In this case, the polymeric  
product is deposited in the lumen of the cavity.

Often times, the hollow or tubular geometry of  
20 organs has functional significance such as in the  
facilitation of fluid or gas transport (blood, urine,  
lymph, oxygen or respiratory gases) or cellular  
containment (ova, sperm). Disease processes may  
affect these organs or their components by encroaching  
25 upon, obstructing or otherwise reducing the cross-  
sectional area of the hollow or tubular elements.  
Additionally, other disease processes may violate the  
native boundaries of the hollow organ and thereby  
affect its barrier function and/or containment  
30 ability. The ability of the organ or structure to  
properly function is then severely compromised. A  
good example of this phenomena can be seen by  
reference to the coronary arteries.

Coronary arteries, or arteries of the heart,  
35 perfuse the actual cardiac muscle with arterial blood.  
They also provide essential nutrients and allow for

removal of metabolic wastes and for gas exchange. These arteries are subject to relentless service demands for continuous blood flow throughout the life of the patient.

5       Despite their critical life supporting function, coronary arteries are often subject to attack through several disease processes, the most notable being atherosclerosis or hardening of the arteries. Throughout the life of the patient, multiple factors  
10       contribute to the development of microscopic and/or macroscopic vascular lesions known as plaques.

          The development of a plaque lined vessel typically leads to an irregular inner vascular surface with a corresponding reduction of vessel cross-  
15       sectional area. The progressive reduction in cross-sectional area compromises flow through the vessel. For example, the effect on the coronary arteries, is a reduction in blood flow to the cardiac muscle. This reduction in blood flow, with corresponding reduction  
20       in nutrient and oxygen supply, often results in clinical angina, unstable angina or myocardial infarction (heart attack) and death. The clinical consequences of the above process and its overall importance are seen in that atherosclerotic coronary  
25       artery disease represents the leading cause of death in the United States today.

          Historically, the treatment of advanced atherosclerotic coronary artery disease i.e. beyond that amenable to therapy via medication alone, involved  
30       cardio-thoracic surgery in the form of coronary artery bypass grafting (CABG). The patient is placed on cardio-pulmonary bypass and the heart muscle is temporarily stopped. Repairs are then surgically affected on the heart in the form of detour conduit  
35       grafted vessels providing blood flow around obstructions. While CABG has been perfected to be quite effective it carries with it inherent surgical risk and requires a several week, often painful

recouperation period. In the United States alone approximately 150-200 thousand people are subjected to open heart surgery annually.

In 1977 a major advance in the treatment of  
5 atherosclerotic coronary artery disease occurred with the introduction of a technique known as Percutaneous Transluminal Coronary Angioplasty (PTCA). PTCA involves the retrograde introduction, from an artery in the arm or leg, up to the area of vessel occlusion,  
10 of a catheter with a small dilating balloon at its tip. The catheter is snaked through the arteries via direct fluoroscopic guidance and passed across the luminal narrowing of the vessel. Once in place, the catheter balloon is inflated to several atmospheres of  
15 pressure. This results in "cracking", "plastic" or otherwise mechanical deformation of the lesion or vessel with a subsequent increase in the cross-sectional area. This in turn reduces obstruction, and trans-lesional pressure gradients and increases blood  
20 flow.

PTCA is an extremely effective treatment with a relatively low morbidity and is rapidly becoming a primary therapy in the treatment of atherosclerotic coronary disease throughout the United States and the  
25 world. By way of example, since its introduction in 1977, the number of PTCA cases now exceeds 150,000 per annum in the United States and, for the first time in 1987, surpassed the number of bypass operations performed. Moreover, as a result of PTCA, emergency  
30 coronary artery bypass surgery is required in less than four percent of patients. Typically, atherosclerosis is a diffuse arterial disease process exhibiting simultaneous patchy involvement in several coronary arteries. Patients with this type of  
35 widespread coronary involvement, while previously not considered candidates for angioplasty, are now being treated due to technical advances and increased clinical experience.

Despite the major therapeutic advance in the treatment of coronary artery disease which PTCA represents, its success has been hampered by the development of vessel renarrowing or reclosure post dilation. During a period of hours or days post procedure, significant total vessel reclosure may develop in up to 10% of cases. This is referred to as "abrupt reclosure". However, the more common and major limitation of PTCA, is the development of progressive reversion of the vessel to its closed condition, negating any gains achieved from the procedure.

This more gradual renarrowing process is referred to as "restenosis." Post-PTCA follow-up studies report a 10-50% incidence (averaging approximately 30%) of restenosis in cases of initially successful angioplasty. Studies of the time course of restenosis have shown that it is typically an early phenomenon, occurring almost exclusively within the six months following an angioplasty procedure. Beyond this six-month period, the incidence of restenosis is quite rare. Despite recent pharmacologic and procedural advances, little success has been achieved in preventing either abrupt reclosure or restenosis post-angioplasty.

Restenosis has become even more significant with the increasing use of multi-vessel PTCA to treat complex coronary artery disease. Studies of restenosis in cases of multi-vessel PTCA reveal that after multi-lesion dilatation, the risk of developing at least one recurrent coronary lesion range from 26% to 54% and appears to be greater than that reported for single vessel PTCA. Moreover, the incidence of restenosis increases in parallel with the severity of the pre-angioplasty vessel narrowing. This is significant in light of the growing use of PTCA to treat increasingly complex multi-vessel coronary artery disease.

The 30% overall average restenosis rate has significant costs including patient morbidity and risks as well as medical economic costs in terms of follow-up medical care, repeat hospitalization and recurrent catheterization and angioplasty procedures. Most significantly, prior to recent developments, recurrent restenosis following multiple repeat angioplasty attempts could only be rectified through cardiac surgery with the inherent risks noted above.

10 In 1987 a mechanical approach to human coronary artery restenosis was introduced by Swiss investigators referred to as, "Intracoronary Stenting". An intracoronary stent is a tubular device made of fine wire mesh, typically stainless steel.

15 The Swiss investigators utilized a stent of the Wallsten design as disclosed and claimed in U.S. Patent No. 4,655,771. The device can be configured in such a manner as to be of low cross-sectional area. In this "low profile" condition the mesh is placed in

20 or on a catheter similar to those used for PTCA. The stent is then positioned at the site of the vascular region to be treated. Once in position, the wire mesh stent is released and allowed to expand to its desired cross-sectional area generally corresponding to the

25 internal diameter of the vessel. Similar solid stents are also disclosed in U.S. Patent No. 3,868,956 to Alfidi et al.

The metal stent functions as a permanent intra-vascular scaffold. By virtue of its material

30 properties, the metal stent provides structural stability and direct mechanical support to the vascular wall. Stents of the Wallsten design are self-expanding due to their helical "spring" geometry. Recently, U.S. investigators introduced slotted steel

35 tubes and extended spring designs. These are deployed through application of direct radial mechanical pressure conveyed by a balloon at the catheter tip. Such a device and procedure are claimed in U.S. Patent

No. 4,733,665 to Palmaz. Despite the significant limitations and potentially serious complications discussed below, this type of stenting has been successful with an almost 100% acute patency rate and a marked reduction in the restenosis rate.

The complications associated with permanent implants such as the Palmaz device result from both the choice of material, i.e., metal or stainless steel, as well as the inherent design deficiencies in the stenting devices. The major limitation lies in the permanent placement of a non-retrievable, non-degradable, foreign body in a vessel to combat restenosis which is predominately limited to the six-month time period post-angioplasty. There are inherent, significant risks common to all permanent implant devices. Moreover, recent studies have revealed that atrophy of the media, the middle arterial layer of a vessel, may occur as a specific complication associated with metal stenting due to the continuous lateral expansile forces exerted after implantation.

These problems are even more acute in the placement of a permanent metallic foreign body in the vascular tree associated with the cardiac muscle. Coronary arteries are subjected to the most extreme service demands requiring continuous unobstructed patency with unimpeded flow throughout the life of the patient. Failure in this system will lead to myocardial infarction (heart attack) and death. In addition, the torsional and other multi-directional stresses encountered in the heart due to its continuous oscillatory/cyclic motion further amplifies the risks associated with a permanent, stiff metallic intra-arterial implant in the coronary bed.

It has been observed that, on occasion, recurrent intravascular narrowing has occurred post-stent placement in vessels during a period of several weeks to months. Typically, this occurs "peri-stent", i.e.,



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immediately up or down stream from the stent. It has been suggested that this may relate to the significantly different compliances of the vessel and the stent, sometimes referred to as "compliance mismatch". Aside from changes in compliance another important mechanism leading to luminal narrowing above and below the stent may be the changes in shear forces and fluid flows encountered across the sharp transitions of the stent-vessel interface. Further supporting evidence has resulted from studies of vascular grafts which reveal a higher incidence of thrombosis and eventual luminal closure also associated with significant compliance mismatch.

To date known stent designs, i.e. tubular, wire helical or spring, scaffold design have largely been designed empirically without consideration or measurement of their radial stiffness. Recent studies measuring the relative radial compressive stiffness of known wire stents, as compared to physiologically pressurized arteries, have found them to be much stiffer than the actual biologic tissue. These studies lend support to the concept of poor mechanical biocompatibility of currently available stents.

Conventional metal stenting is severely limited since it is device dependent and necessitates a myriad of individual stents as well as multiple deployment catheters of varying lengths and sizes to accommodate individual applications. Additionally, metal stents provide a relatively rigid nonflexible structural support which is not amenable to a wide variety of endoluminal geometries, complex surfaces, luminal bends, curves or bifurcations.

These identified risks and limitations of metal stents have severely limited their utility in coronary artery applications. As of 1988, a partial self-imposed moratorium exists in the use of helical metal stents to treat human coronary artery diseases. Presently in the United States, a spring-like wire

coil stent has been approved only for short term use as an emergency device for patients with irreparably closed coronary arteries following failed PTCA while in transit to emergency bypass surgery. An  
5 alternative to the use of stents has now been found which has broad applications beyond use in coronary artery applications for keeping hollow organs open and in good health.

#### Summary of the Invention

10 The present invention provides a solution to the problem of restenosis following angioplasty, without introducing the problems associated with metal stents. Specifically, the invention provides a novel method for endoluminal paving and sealing (PEPS) which  
15 involves application of a polymeric material to the interior surface of the involved blood vessel. In accordance with this method, a polymeric material, either in the form of a monomer or prepolymer solution or as an at least partially pre-formed polymeric  
20 product, is introduced into the lumen of the blood vessel and positioned at the point of the original stenosis. The polymeric product is then reconfigured to conform to and maintain intimate contact with the interior surface of the blood vessel such that a  
25 paving and sealing coating is achieved.

The PEPS approach is not limited to use in connection with restenosis, however, and can also be effectively employed in any hollow organ to provide local structural support, a smooth surface, improved  
30 flow and sealing of lesions. In addition, the polymeric paving and sealing material may incorporate therapeutic agents such as drugs, drug producing cells, cell regeneration factors or even progenitor cells of the same type as the involved organ or  
35 histologically different to accelerate healing processes. Such materials with incorporated therapeutic agents may be effectively used to coat or

plug surgically or traumatically formed lumens in normally solid organs as well as the native or disease generated lumens of hollow or tubular organs.

For use in these applications, the present  
5 invention provides at least partially preformed polymeric products. These products may have any of a variety of physical shapes and sizes in accordance with the particular application. The invention also provides apparatus specially adapted for the  
10 positioning of the polymeric material, these including partially pre-formed polymeric products, at the interior surface of an organ and for the subsequent chemical or physical reconfiguration of the polymeric material such that it assumes a desired molded or  
15 customized final configuration.

#### Brief Description of the Drawing

Fig. 1 shows an amorphous geometry of the PEPS polymer coating before and after deployment;

Fig. 2 shows a stellate geometry of the PEPS  
20 polymer coating before and after deployment;

Fig. 3 shows a linear feathered polymer strip applied to "one" wall before and after deployment;

Fig. 4 shows a large patch of sprayed on polymer material before and after deployment;

25 Fig. 5 shows a porous tubular form geometry before and after deployment;

Fig. 6 shows a spot geometry of the PEPS process before and after deployment;

Fig. 7 shows a spiral form application of the  
30 PEPS process before and after deployment;

Fig. 8 shows an arcuate (radial, arc) patch geometry of the PEPS polymer before and after deployment;

Fig. 9 shows a process for using PEPS to treat an  
35 artificially created tissue lumen;

Fig. 10 shows two lumen catheters according to the invention;

Fig. 11 shows surface contours of expansile members useful in catheters according to the invention;

Fig. 12 shows three catheters according to the invention;

Fig. 13 shows four lumen catheters according to the invention;

Fig. 14 shows five lumen catheters according to the invention;

Fig. 15 shows six lumen catheters according to the invention;

Fig. 16 shows seven lumen catheters according to the invention;

Fig. 17 shows a distal occlusion catheter and a polymer delivery catheter in a vessel;

Fig. 18 shows in cross-section a polymeric sleeve before insertion in a blood vessel; the sleeve after insertion in the vessel and after expansion;

Fig. 19 is a cross-sectional comparison of an initial polymeric sleeve and an expanded polymeric sleeve.

Fig. 20 shows enmeshed discontinuous polymeric material arrayed on a catheter with a retractable sheath; and

Fig. 21 shows variations in apertures for polymer delivery to a tissue lumen.

#### Detailed Description of the Invention

In general, PEPS involves the introduction of a polymeric material into a selected location within a lumen in tissue, i.e. an organ, an organ component or cavernous component of an organism, and the subsequent reconfiguration of the polymeric material to form a sealing in intimate and conforming contact with or paving the interior surface. As used herein, the term "sealing" or "seal" means a coating of sufficiently low porosity that the coating serves a barrier function. The term "paving" refers to coatings which

are porous or perforated. By appropriate selection of the polymeric material employed and of the configuration of the coating or paving, PEPS provides a unique customizable process, which can be utilized  
5 as a given biological or clinical situation dictates.

The basic requirements for the polymeric material to be used in the PEPS process are biocompatibility and the capacity to be chemically or physically reconfigured under conditions which can be achieved in  
10 vivo. Such reconfiguration conditions may involve heating, cooling, mechanical deformation, e.g., stretching, or chemical reactions such as polymerization or crosslinking.

Suitable polymeric materials for use in the  
15 invention include polymers and copolymers of carboxylic acids such as glycolic acid and lactic acid, polyurethanes, polyesters such as poly(ethylene terephthalate), polyamides such as nylon, polyacrylonitriles, polyphosphazines, polylactones  
20 such as polycaprolactone, and polyanhydrides such as poly[bis(p-carboxyphenoxy)propane anhydride] and other polymers or copolymers such as polyethylene, polyvinyl chloride and ethylene vinyl acetate.

Other bioabsorbable polymers could also be used  
25 either singly or in combination, or such as homopolymers and copolymers of delta-valerolactone, and p-dioxanone as well as their copolymers with caprolactone. Further, such polymers can be cross-linked with bis-caprolactone.

30 Preferably PEPS utilizes biodegradable polymers, with specific degradation characteristics to provide sufficient lifespan for the particular application. As noted above, a six month lifespan is probably sufficient for use in preventing restenosis; shorter  
35 or longer periods may be appropriate for other therapeutic applications.

Polycaprolactone as disclosed and claimed in U.S. Patent No. 4,702,917 to Schindler, incorporated herein

by reference, is a highly suitable bioabsorbable polymer for use in the PEPS process, particularly for prevention of restenosis. Polycaprolactone possesses adequate mechanical strength being mostly crystalline even under quenching conditions. Despite its structural stability, polycaprolactone is much less rigid than the metals used in traditional stenting. This minimizes the risk of acute vessel wall damage from sharp or rough edges. Furthermore, once polycaprolactone has been deployed its crystalline structure will maintain a constant outside diameter. This eliminates the risks often associated with known helical or spring metal stents which after being expanded in vivo have a tendency to further expand exerting increasing pressure on the vessel wall.

The rate of bioabsorption of polycaprolactone is ideal for this application. The degradation process of this polymer has been well characterized with the primary degradation product being nonparticulate, nontoxic, 6-hydroxy hexanoic acid of low acidity. The time of biodegradation of polycaprolactone can be adjusted through the addition of various copolymers.

Polycaprolactone is a preferred polymer for use in the PEPS process because it has attained favorable clinical acceptability and is in the advanced stages of FDA approval. Polycaprolactone has a crystalline melting point of 60°C and can be deployed in vivo via a myriad of techniques which facilitate transient heating and varying degrees of mechanical deformation or application as dictated by individual situations. This differs markedly from other bioabsorbable polymers such as polyglycolide and polylactide which melt at much higher temperatures of 180°C and pose increased technical constraints as far as the delivery system affording polymer sculpting without deleterious tissue exposure to excessive temperatures or mechanical forces.

Polyanhydrides have been described for use as drug carrier matrices by Leong et al., J. Biomed. Mat. Res. 19, 941-955 (1985). These materials frequently have fairly low glass transition temperatures, in some cases near normal body temperature, which makes them mechanically deformable with only a minimum of localized heating. Furthermore, they offer erosion times varying from several months to several years depending on particular polymer selected.

- 10 The polymeric materials may be applied in custom designs, with varying thicknesses, lengths, and three-dimensional geometries (e.g. spot, stellate, linear, cylindrical, arcuate, spiral) to achieve varying finished geometries as depicted in Figs. 1-8.
- 15 Further, PEPS may be used to apply polymer to the inner surfaces of hollow, cavernous, or tubular biological structures (whether natural or artificially formed) in either single or multiple polymer layer configurations. PEPS may also be used, where
- 20 appropriate, to occlude a tissue lumen completely.

- The polymeric material used in PEPS can be combined with a variety of therapeutic agents for on-site delivery. Examples for use in coronary artery applications are anti-thrombotic agents, e.g.,
- 25 prostacyclin and salicylates, thrombolytic agents e.g. streptokinase, urokinase, tissue plasminogen activator (TPA) and anisoylated plasminogen-streptokinase activator complex (APSAC), vasodilating agents i.e. nitrates, calcium channel blocking drugs, anti-
- 30 proliferative agents i.e. colchicine and alkylating agents, intercalating agents, growth modulating factors such as interleukins, transformation growth factor  $\beta$  and congeners of platelet derived growth factor, monoclonal antibodies directed against growth
- 35 factors, anti-inflammatory agents, both steroidal and non-steroidal, and other agents which may modulate vessel tone, function, arteriosclerosis, and the healing response to vessel or organ injury post

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intervention. In applications where multiple polymer layers are used different pharmacological agents could be used in different polymer layers. Moreover, PEPS may be used to effect pharmaceutical delivery focally within the vessel wall, i.e. media.

The polymeric material in accordance with the invention may also have incorporated in it living cells to serve any of several purposes. For examples, the cells may be selected, or indeed designed using principles of recombinant DNA technology, to produce specific agents such as growth factors. In such a way, a continuously regenerating supply of a therapeutic agent may be provided without concerns for stability, initial overdosing and the like.

Cells incorporated in the polymeric material may also be progenitor cells corresponding to the type of tissue in the lumen treated or other cells providing therapeutic advantage. For example, liver cells might be implanted in the polymeric material within a lumen created in the liver of a patient to facilitate regeneration and closure of that lumen. This might be an appropriate therapy in the case where scar tissue or other diseased, e.g. cirrhosis, fibrosis, cystic disease or malignancy, or non-functional tissue segment has formed in the liver or other organ and must be removed. The process of carrying out such treatment, shown schematically in Fig. 9, involves first inserting a catheter 91 into a lumen 92 within a diseased organ segment 93. The lumen 92 can be a native vessel, or it can be a man-made lumen, for example a cavity produced by a laser. The catheter 91 is used to introduce a polymeric plug 94 into the lumen 92. The catheter is then removed, leaving the plug 94 in place to act as a focus for new growth stemming from cells implanted along with the polymeric plug 94. If the desire is for a more tubular structure, the plug 94 can be appropriately reconfigured.



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Optional additions to the polymeric material such as barium, iodine or tantalum salts for X-ray radio-opacity allow visualization and monitoring of the coating.

- 5       The technique of PEPS preferably involves the percutaneous application of a polymeric material, preferably a biodegradable polymer such as polycaprolactone, either alone or mixed with other biodegradable polymeric materials, which may
- 10       optionally contain various pharmaceutical agents for controlled sustained release of the pharmaceutical or for selective soluble factor adsorption and trapping. The polymeric material is typically applied to the inside of an organ surface employing combined thermal
- 15       and mechanical means to manipulate the polymeric material. Although capable of being used during surgery, PEPS will generally be applied without the need for a surgical procedure using some type of catheter, for example novel modifications of the known
- 20       catheter technology described above for (PTCA). PEPS is preferably applied using a single catheter with multiple balloons and lumens. The catheter should be of relatively low cross-sectional area. Typically a long thin tubular catheter manipulated using
- 25       fluoroscopic guidance can access deep into the interior of organ or vascular areas.

- The polymer may be deployed in the interior of the vessel or organ from the surface or tip of the catheter. Alternatively, the polymer could be
- 30       positioned on a balloon such as that of a standard angioplasty balloon catheter. Additionally, the polymer could be applied by spraying, extruding or otherwise internally delivering the polymer via a long flexible tubular device consisting of as many lumens
- 35       as a particular application may dictate.

      The simplest PEPS coating is a continuous coating over a designated portion of a tissue lumen. Such a coating can be applied with a simple two lumen

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catheter such as those shown in Fig. 10. Looking first to Fig. 10a, a suitable catheter is formed from a tubular body 100 having a proximal end 101 and a distal end 102. The interior of the tubular body 100 is divided into two conduits 103 and 104 which extend from the proximal end 101 to apertures 105 and 106 in the tubular body. (Figs. 10b and 10c) Conduits 103 and 104 thus connect apertures 105 and 106 with the proximal end 101 of the tubular body 100 to allow fluid flow therebetween. The proximal ends of conduits 103 and 104 are preferably equipped with connectors 108 which allow connection with fluid supplies. Pressure connectors such as Luer® locks are suitable.

The catheter may also include markers 109 in one or more locations to aid in locating the catheter. These markers can be, for example, fluoroscopic radio-opaque bands affixed to the tubular body 100 by heat sealing.

The catheter shown in Figs. 10b and 10c has an expansile member in the form of an inflatable balloon 107 disposed over the distal aperture 105. In use, an at least a partially preformed polymeric layer or partial layer is positioned over the balloon 107 and the catheter is inserted into an appropriate position in the tissue lumen. Fluid flow through conduit 103 will cause the balloon 107 to inflate, stretch and deform the polymer layer until it comes into contact with the walls of the tissue lumen. The other aperture 105 and conduit 103 are used to control the reconfiguration of the polymeric sleeve, for example by supplying a flow of heated liquid to soften the sleeve and render it more readily stretchable or to stimulate polymerization of a partially polymerized sleeve.

Variations on this basic two lumen catheter can be made, examples of which are shown in Figs. 10d and 10e. For example Fig. 10d has a shapable wire affixed

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to the tip of the catheter to aid in insertion and a traumatic and directed passage through the organism, i.e. to act as a guide wire. In Fig. 10e, the expansile member is incorporated as part of the tubular body as a continuous element, preferably a unitary element. In this case, the distal tip 107a of the catheter expands in response to fluid flow in conduit 103. Conduit 104 can be formed by bonding in or on the extruded catheter body a piece of the same or different material in a tubular form. This type of design can also be used in more complicated multi-lumen catheters discussed below.

The polymeric material may take the form of a sleeve designed to be readily insertable along with the catheter into the tissue lumen, and then to be deployed onto the wall of the lumen to form the coating. This deployment can be accomplished by inflating a balloon, such as balloon 107 using fluid flow through conduit 103. Inflation of balloon 107 stretches the polymeric sleeve causing it is press against the walls of the tissue lumen and acquire a shape corresponding to the lumen wall. This shape is then fixed, and the catheter removed leaving behind a polymeric paving or seal on the lumen wall.

The process of fixing the shape of the polymeric material can be accomplished in several ways, depending on the character of the original polymeric material. For example, a partially polymerized material can be expanded using the balloon after which the conditions are adjusted such that polymerization can be completed, e.g., by increasing the local temperature or providing UV radiation through an optical fiber. A temperature increase might also be used to soften a fully polymerized sleeve to allow expansion and facile reconfiguration and local molding, after which it would "freeze" in the expanded position when the heat source is removed. Of course, if the polymeric sleeve is a plastic material which

will permanently deform upon stretching (e.g., polyethylene, polyethylene terephthalate, nylon or polyvinyl chloride), no special fixation procedure is required.

- 5 As depicted in Fig. 10b, local heating can be provided by a flow of heated liquid directly into the tissue lumen. Thermal control can also be provided, however, using a fluid flow through or into the expansile member or using a "leaky" partially  
10 perforated balloon such that temperature control fluid passes through the expansile member, or using electrical resistive heating using a wire running along the length of the catheter body in contact with resistive heating elements. This type of heating  
15 element can make use of DC or radiofrequency (RF) current or external RF or microwave radiation. Other methods of achieving temperature control can also be used, including laser heating using an internal optical fiber (naked or lensed) or thermonuclear  
20 elements.

- In addition to the smooth shape shown in Fig. 10, the balloon used to configure the polymer can have other surface shapes for formation of the coatings to provide specific polymeric deployment patterns. For  
25 example, the balloon may be a globular shape intended for deployment from the tip of a catheter device. (Fig. 11a) Such an arrangement would be preferred when the paving operation is being carried out in a cavity as opposed to a tubular organ. The balloon  
30 might also be thickened at the ends (Fig. 11c) or substantially helical (Fig. 11d) providing a variation in coating thickness along the length of the paved or sealed area. Such a configuration might prove advantageous in the case where additional structural  
35 support is desired and to provide a tapered edge to minimize flow disruption. Variations in coating thickness which provide ribs running the length of the tissue lumen might be achieved using a stellate

balloon (Fig. 11e). This type of polymer coating would be useful in the case where additional structural support is desirable combined with more continuous flow properties. In addition balloon shape  
5 may facilitate insertion in some cases.

Variations in the ultimate configuration of the PEPS coating can also be achieved by using more complex deployments of the polymer on the expansile member. For example, the polymer can be in the form  
10 of a perforated tubular sleeve, a helical sleeve or in the form of discontinuous members of various shapes. These may be affixed to the expansile member directly, for example with an adhesive or by suction through perforations and the like, or to an overcoating such  
15 as dissolvable gauze-like or paper sheath (i.e. spun saccharide) or held in place by or a retractable porous sheath which will be removed with the catheter after application.

For example, Fig. 20(a) shows an array of polymer  
20 dots. These dots are enmeshed in a dissolvable mesh substrate Fig. 20(b) which in turn is wrapped around the expansile member 107 of a catheter according to the invention (Fig. 20c). An exemplary two lumen catheter is shown in Fig. 20d (numbered as in Fig.  
25 10b) where a retractable sheath 205 surrounds the polymer dots 206 for insertion. When the catheter reaches the application site, the sheath 205 is retracted (Fig. 20e) and the balloon 107 expanded.

It will be recognized, that the catheter depicted  
30 in Fig. 10 represents a minimalist approach to PEPS catheter design, and that additional lumens may be included within the catheter body to provide conduits for inflation of positioning balloons, optical fibers, additional polymer molding balloons, temperature  
35 control means, and passage of steering or guide wires or other diagnostic devices, e.g. ultrasound catheter, or therapeutic devices such as atherectomy catheter or other lesion modifying device. For example, three

lumen catheters (Fig. 12), four lumen catheters (Fig. 13), five lumen catheters (Fig. 14), six lumen catheters (Fig. 15) and seven lumen catheters (Fig. 16) might be employed. A retractable sheath may also be provided which extends over the polymer during insertion to prevent premature separation of the polymer from the catheter. In addition, catheters may have telescoping sections such that the distance between the occluding balloons can be varied.

Looking for example at the six lumen catheters in Fig. 15b, two positioning balloons 150 and 151, both connected to conduit 152. Positioning balloons 150 and 151 serve to fix the position of the tubular body 100 within a tissue lumen and isolate the portion of the tissue lumen between them where the PEPS coating will be applied. Expansile member 153 is provided with circulating flow via conduits 154 and 155. This can be used to provide temperature control to the isolated portion of the tissue lumen, as well as acting to configure the polymeric coating formed by expanding a polymeric sleeve and other deployed form fitted over expansile member 153. In the catheter shown in Fig. 15b, a temperature control solution or a therapeutic solution is advantageous provided through conduit 156, with conduit 157 acting as a drain line (or vice versa) to allow flow of fluid through the isolated portion of the tissue lumen ("superfusion"). Such a drain line is not required, however, and a simple infusion catheter could omit one of the conduits 156 or 157 as in the five lumen designs of Fig. 14. The sixth conduit 158 is also optional, but can be advantageously used for guide wires, diagnostic or therapeutic device passage, or distal fluid perfusion. If conduit 158 has an aperture proximal to balloon 151, it can be used as a by-pass conduit for passive perfusion during occlusion.

The incorporation in the catheter of positioning balloons which occlude a section of the tissue lumen

makes it possible to utilize solutions of monomers or prepolymers and form the coating in situ. Looking for example at four lumen catheters shown in Fig. 13b, an isolation zone is created by inflating balloons 131 and 132 so that they press against the tissue lumen. While expansile member 133 could be used to deform a polymeric sleeve or other deployment form, it can also be used to define the size and environmental conditions (e.g. temperature) of the lumen region.

Application of the polymeric material may be accomplished by extruding a solution of monomers or prepolymers through the aperture 134 to coat or fill the tissue lumen. The formation of a polymer coating can be controlled by introducing crosslinking agents or polymerization catalysts together with the monomer or prepolymer solution and then altering the conditions such that polymerization occurs. Thus, a flow of heated fluid into expansile member 133 can increase the local temperature to a level sufficient to induce or accelerate polymerization.

Alternatively, the monomer/prepolymer solution might be introduced cold, with metabolic temperature being sufficient to induce polymerization. The other lumen 135 acts as a drain line in superfusion applications.

The polymeric material can be introduced to the tissue lumen through a simple aperture in the side of the tube as shown in Fig. 21a, or through a raised aperture (Fig. 21b). A shaped nozzle which is extendable away from the surface of the tubular body (Fig. 21e) can also be used. The material can be extruded through, or it can be subjected to flow restriction to yield a spray application. This flow restriction can be adjustable to control the spray. In addition, localized acceleration at the tip of the nozzle can be used, for example, via a piezoelectric element to provide sprayed application.

The catheters bodies for use in this invention can be made of any known material, including metals,

e.g. steel, and thermoplastic polymers. Occluding balloons may be made from compliant materials such as latex or silicone, or non-compliant materials such as polyethyleneterephthalate (PET). The expansile member  
5 is preferably made from non-compliant materials such as PET, PVC, polyethylene or nylon. The expansile member may optionally be coated with materials such as silicones, polytetra-fluoroethylene (PTFE), hydrophilic materials like hydrated hydrogels and  
10 other lubricious materials to aid in separation of the polymer coating.

In addition to arteries, i.e. coronary, femoro-iliac, carotid and vertebro-basilar, the PEPS process may be utilized for other applications such as paving  
15 the interior of veins, ureters, urethrae, bronchi, biliary and pancreatic duct systems, the gut, eye and spermatic and fallopian tubes. The sealing and paving of the PEPS process can also be used in other direct clinical applications even at the coronary level.

20 These include but are not limited to the treatment of abrupt vessel reclosure post PCTA, the "patching" of significant vessel dissection, the sealing of vessel wall "flaps", i.e. secondary to catheter injury or spontaneously occurring, the sealing of aneurysmal  
25 coronary dilations associated with various arteritides. Further, PEPS provides intra-operative uses such as sealing of vessel anastomoses during coronary artery bypass grafting and the provision of a bandaged smooth polymer surface post endarterectomy.

30 The unique pharmaceutical delivery function of the PEPS process may be readily combined with "customizable" deployment geometry capabilities to accommodate the interior of a myriad of complex organ or vessel surfaces. Most importantly, this customized  
35 geometry can be made from structurally stable yet biodegradable polymers. The ability to tailor the external shape of the deployed polymer through melted polymer flow into uneven surface interstices, while



maintaining a smooth interior surface with good flow characteristics, will facilitate better structural support for a variety of applications including eccentric coronary lesions which by virtue of their geometry are not well bridged with conventional metal stents.

As noted above, the polymer substrate used in PEPS may be fashioned, for example, out of extruded tubes of polycaprolactone and/or copolymers. The initial predeployment design and size of the polymer sleeve will be dictated by the specific application based upon the final deployed physical, physiological and pharmacological properties desired.

For coronary artery application, predeployment tubes of about 10 to 20 mm in length and about 1 to 2 mm in diameter would be useful. The initial wall thickness of the resulting in vivo polymer layer may be varied depending upon the nature of the particular application. In general coating procedures require polymer layers of about 0.005mm to 0.50mm while layers which are designed to give structural support can vary from 0.05mm to 5.0mm.

The polymer tube walls may be processed prior to insertion with either laser or chemical etching, pitting, slitting or perforation depending upon the application. In addition, the shape of any micro (10nm to 1  $\mu$ m) or macro (>1  $\mu$ m up to about 15  $\mu$ m) perforation may be further geometrically modified to provide various surface areas on the inner versus outer seal surface. The surfaces of the predeployed polymer may be further modified with bound, coated, or otherwise applied agents, i.e., cyanoacrylates or biological adhesives such as those derived from fungal spores, the sea mussel or autologous fibrinogen adhesive derived from blood.

For PEPS applications involving the coronary arteries, the polymer tubes (if in an initial tubular configuration), should preferably have perforations or

pores, of a size dictated by the particular application. This will ensure a symmetric expansion of the encasing polymeric sealant. By using a fragmented tubular polymer surface with corresponding  
5 expansions along predicted perforations (i.e., the slots) a significant mechanical stability is provided. In addition, this minimizes the amount of foreign material placed within the vessel.

Depending upon the polymer and pharmaceutical  
10 combination and the configuration, PEPS may be used to coat or bandage the organ inner surface with a thin adhesive partitioning polymer film or layer of about 0.005mm to 0.50mm. Biodegradable polymers thus applied to an internal organ or vessel surface will  
15 act as an adherent film "bandage." This improved surface, with desirable rheologic and adherence properties, facilitates improved fluid or gas transport in and through the body or lumen of the vessel or hollow organ structure and acts to reinstate  
20 violated native surfaces and boundaries.

The ultimate in vivo deployed geometry of the polymer dictates the final function of the polymer coating. The thinner applications allow the polymer film to function as a coating, sealant and/or  
25 partitioning barrier, bandage, and drug depot. Complex internal applications of thicker layers of polymer, such as intra-vessel or intra-luminal applications, may actually provide increased structural support and depending on the amount of  
30 polymer used in the layer may actually serve in a mechanical role to maintain vessel or organ potency.

For example, lesions which are comprised mostly of fibromuscular components have a high degree of visco-elastic recoil. These lesions would require  
35 using the PEP process to apply an intraluminal coating of greater thickness and extent so as to impart more structural stability thereby resisting vessel radial compressive forces. The PEPS process in this way

provides structural stability and is generally applicable for the maintenance of the intraluminary geometry of all tubular biological organs or substructure. It may be used in this way following the therapeutic return of normal architecture associated with either balloon dilation (PTCA), atherectomy, lesion spark, thermal or other mechanical erosion, "G-lazing", welding or laser recanalization.

An important feature of the PEPS technique is the ability to customize the application of the polymer to the internal surface of a vessel or organ as dictated by the particular application. This results in a variety of possible geometries of polymer as well as a variety of forms. These multi-geometry, multi-form polymer structures may be adjusted to correspond to particular functions. (Figs. 1-8)

With particular reference to Figs. 1-8 the PEPS process may be affectuated so that the focal application of polymer to the vessel or organ results in either an amorphous geometry, Fig. 1, stellate geometry, Fig. 2, or spot geometry, Fig. 6. Additional geometries could include a linear feathered polymer strip applied to a particular area of the vessel wall as shown in Fig. 3. Fig. 4 shows a large patch of polymer which can be sprayed on using a variety of known techniques. Another form of the PEPS application to be utilized in instances, e.g., where structural stability need be imparted to the vessel would be the porous tubular form shown in Fig. 5. Other types of PEPS applications which would impart structural stability to the vessel would be the spiral form application shown in Fig. 7, or the arcuate (radial, arc) patch as shown in Fig. 8.

Conversely, in cases where the severely denuded lesions have irregular surfaces with less fibromuscular components, the PEPS process can be used to provide only a thin polymer film to act as a bandage.

The PEPS' process is significantly different and is conceptually an advance beyond stents and stenting in achieving vessel patency. Stents have been designed with the underlying primary function of providing a relatively stiff structural support to resist post PTCA, vessel reclosure caused by the vessel's spring-like characteristics. It has been increasingly demonstrated that cellular and biochemical mechanisms as opposed to physical "spring-like" coils, are of a much greater significance in preventing vessel reclosure and PEPS addresses these mechanisms.

The specific object and features of the PEPS process are best understood by way of illustration with reference to the following examples and figures.

Example 1

The invention may be readily understood through a description of an experiment performed in vitro using a mock blood vessel made from transparent plastic tubing using a heat-balloon type deployment catheter reference to Fig. 17.

The balloon delivery catheter 170 is first positioned in the vessel 171 at the area of the occlusion. Before insertion, a polycaprolactone polymer sleeve 172 containing additives, e.g. to aid X-ray radio-opacity, for drug delivery or to promote surface adhesion, is placed in a low profile condition surrounding a balloon at the distal end of the delivery catheter 170. The delivery catheter with the polycaprolactone tube is then inserted balloon end first into the vessel 171 and manipulated into position, i.e., the area of the vessel to be treated.

A separate occlusion catheter 173 is employed to restrict "blood" flow through the vessel. The distal end of the occlusion catheter 173 is inflated to create a stagnant column of "blood" in the vessel around the balloon delivery catheter and polycaprolactone tube. Saline solution at about 60-

80°C is injected through a lumen in the occlusion catheter 173 or the delivery catheter 170 in the case of using a catheter according to the invention into the area surrounding the delivery catheter, balloon and polycaprolactone tube. Once the polycaprolactone tube becomes pliable, the delivery catheter balloon is inflated to push the polycaprolactone sleeve out against the interior wall thereby locally sealing and/or paving the vessel.

10 The polycaprolactone expands and/or flows, conforming to the inner surface of the vessel, flowing into and filling in surface irregularities thereby creating a "tailored" fit. Further, the deployed interior surface of the PEPS polymer is smooth  
15 providing an increased vessel (lumen) cross-section diameter and a rheologically advantageous surface with improved blood flow. Upon removal of heated saline solution the polymer recrystallizes to provide a paved surface of the vessel wall interior.

20 The deployment catheter balloon is then deflated leaving the polycaprolactone layer in place. The balloon section of the occlusion catheter is deflated and, blood flow was allowed to return to normal and the deployment catheter was removed leaving the  
25 recrystallized polycaprolactone layer in place within the vessel.

Over the course of time the polycaprolactone seal will become covered with a proteinaceous biologic thin film coat. Depending upon the exact seal chemical  
30 composition, the polymer will then biodegrade, at a predetermined rate and "dissolve" into the bloodstream or be absorbed into the vessel wall. While in intimate contact with the vessel wall, pharmacological agents if embedded or absorbed in the polycaprolactone  
35 will have a "downstream" effect if released slowly into the bloodstream or may have a local effect on the blood vessel wall, thereby facilitating healing of the angioplasty site, controlling or reducing exuberant

medial smooth muscle cell proliferation, promoting effective lesion endothelialization and reducing lesion thrombogenicity.

Example 2

5 Polycaprolactone in an initial macroporous tubular configuration was placed in a low profile form in bovine coronary arteries and canine carotid arteries. In the process of deployment the vessels were purposely overextended and sealed through thermal  
10 and mechanical deformation of the polymer. Fig. 18 shows a cross-section of the polymer tube 180 before insertion in the bovine artery, after insertion in the artery 181, and after expansion 182. The initial polymer tube 180, is smaller in diameter than the  
15 artery 181. After deployment, the thin film of polymer 182 can be seen coating the inner surface of the sealed vessel with the vessel remaining erect. The vessel remained dilated to about 1.5 times the original diameter because of the ability of the  
20 polymer to keep it fixed. Fig. 19 shows a cross-section of the polymer before insertion 190 and removed after insertion and reconfiguration 191 in a canine artery. This figure clearly shows the stretching and thinning of the polymer wall.

25 All polymer sealed vessels remained dilated with a thin layer of macroporous polymer providing a new barrier surface between the vessel lumen and the vessel wall constituents. The unsealed portion of the vessels did not remain dilated.

30 These examples demonstrate that the PEPS process may if desired provide polymer application with a relatively large degree of surface area coverage and an effective polymer barrier shield. As such, the polymer barrier-shield may, if desired, impart  
35 sufficient structural stability to maintain a selected vessel diameter. The selected final vessel diameter at which a vessel is sealed is dictated by the

particular physiological variables and therapeutic goals which confront the PEPS user.

The geometry of the pre and post PEPS application sites may be readily varied. PEPS may be used to  
5 merely coat an existing vessel or organ geometry. Alternatively, the PEPS process may be used to impart structural stability to a vessel or organ the geometry of which was altered prior to the PEPS application. In addition, the PEPS process may itself alter the  
10 geometry of the vessel or organ by shaping the geometry. With reference to Fig. 18 this latter process was used to expand the vessel 181.

A specific and important attribute of the PEPS technique and the polymers which are employed is the  
15 significantly lower degree of compliance mismatch or similarities of stiffness (inverse of compliance) between the vessel and the polymer seal as compared to metal stents. The vessel damage from compliance mismatch discussed above may be eliminated by the PEPS  
20 process utilizing a variety of available polymers. Additionally, compliance mismatch greatly modifies the characteristics of fluid wave transmission along the vessel with resultant change in local flow properties, development of regional change in shear forces and a  
25 subsequent vessel wall hypertrophy which acts to reduce vessel cross-sectional area and reduces blood flow. Further, the substructural elimination of compliance mismatch of the PEPS technique at first minimizes and then, upon dissolution eliminates local  
30 flow abnormalities and up and downstream transition zone hypertrophy associated with metal stenting.

PEPS has the flexibility of being safely and effectively used prophylactically at the time of initial PTCA in selected patients or being  
35 incorporated as part of the original dilation procedure as a second stage prophylactic vessel surface "finishing" process. For example, the invasive cardiologist may apply the PEPS technique on

a wide clinical basis after the first episodes of restenosis. In addition, because the PEPS technique significantly aids in the vascular healing process post intervention, it may be readily used prophyl-  
5 actically after initial angioplasty prior to any incidence of restenosis. This would free the patient from the risks of repeat intracoronary procedure as well as those associated with metal stenting.



Claims

1. A catheter device for application of a polymeric coating to a tissue lumen comprising a flexible tubular body having proximal and distal ends,  
5 which tubular body defines a lumen divided into a plurality of sublumens, each sublumen extending from the proximal end of the tubular body toward the distal end of the tubular body and connecting to at least one aperture in the tubular body  
10 whereby each sublumen forms a conduit for fluid flow between at least one aperture in the tubular body and the proximal end of the tubular body, characterized in that at least one of said sublumens controls delivery of polymeric coating  
15 material to the tissue lumen.
2. A catheter device as in claim 1, characterized in that the catheter includes at least one occluding balloon member disposed around the tubular body in alignment with the aperture of a sublumen,  
20 whereby fluid flow through the sublumen inflates the positioning balloon member.
3. A catheter device according to claims 1-2, characterized in that the device includes two occluding balloons, one disposed toward the  
25 proximal end of the tubular body from the aperture of the sublumen which controls delivery of polymeric coating material, and the other disposed toward the distal end of the tubular body from the aperture of the sublumen which  
30 controls delivery of polymeric coating material, whereby the occluding balloons together act to at least partially occlude a portion of the tissue lumen.

4. A catheter device as in claim 3, characterized in that the sublumen which controls polymer delivery forms a conduit between a vessel containing a monomer or prepolymer solution for the polymer coating and the tissue lumen.  
5
5. A catheter device as in claims 1-3, characterized in that an expansile member is disposed around the tubular body in alignment with the aperture of the sublumen that controls polymer delivery, whereby fluid flow through the sublumen that controls polymer delivery inflates the expansile member.  
10
6. A catheter device as in claim 5, characterized in that a partial or complete layer of polymeric coating material is disposed around the expansile member, whereby expansion of the balloon causes the polymeric material to expand to coat the tissue lumen.  
15
7. A catheter device as in claims 5-6, characterized in that a continuous, perforated or helical polymeric sleeve is disposed around the expansile member, whereby expansion of the balloon causes the polymeric material to expand to coat the tissue lumen.  
20
8. A catheter device as in claims 5-6, characterized in that the polymeric coating material is discontinuous and disposed on a mesh support.  
25
9. A catheter device as in claim 4, characterized in that there is an expansile member disposed around the tubular body in alignment with a further of said apertures and between the positioning balloons, whereby the thickness and surface  
30

configuration of the polymer coating is controlled by inflation of the expansile member.

10. A catheter device as in claims 5-9, characterized in that the expansile member has a contoured surface whereby a molded shape is imparted to the polymeric coating.  
5
11. A catheter device as in claims 1-4 and 9-10, characterized in that the distal aperture of the sublumen controlling polymer deliver is shaped to provide flow acceleration.  
10
12. A catheter device as in claim 11, characterized in that the shaped aperture includes a flow propellant device.
13. A catheter device as in claims 3-12, characterized in that inflation of the two occluding balloons is independently controllable.  
15
14. A catheter device as in claims 1-13, characterized in that the device includes a heating element, whereby the polymeric coating material can be heated prior or subsequent to delivery to the tissue lumen.  
20
15. A catheter device as in claims 1-14, characterized in that the device includes an optical fiber whereby light can be delivered to the polymeric coating material.  
25
16. A catheter device as in claims 1-3, 5-8 and 13-15 characterized in that the device includes a mechanically retractable protective sheath.

17. A catheter device as in claims 1-17,  
characterized in that the polymeric material  
includes one or more therapeutic agents.
- 5 18. An at least partially preformed polymeric product  
for use in endoluminal paving and sealing.
19. A polymeric product as in claim 18, characterized  
in that the polymer employed is partially  
polymerized.
- 10 20. A polymeric product as in claim 19, characterized  
in that further polymerization of the polymeric  
product is thermally activated.
21. A polymeric product as in claim 19, characterized  
in that further polymerization of the polymeric  
product is photoactivated.
- 15 22. A polymeric product as in claims 18-21,  
characterized in that the polymer is  
biodegradable.
- 20 23. A polymeric product as in claim 22, characterized  
in that the product comprises polygluconolactone  
or a polyanhydride.
24. A polymeric product as in claims 18-21,  
characterized in that the product comprises  
polyethylene, polyvinyl chloride or ethylene  
vinyl acetate copolymer.
- 25 25. A polymeric product as in claims 18-24,  
characterized in that the polymer is mechanically  
deformable at temperatures achievable in vivo.

26. A polymeric product as in claims 18-25,  
characterized in that the product comprises one  
or more therapeutic or cell growth agents.
27. A polymeric product as in claims 18-26,  
5 characterized in that the product is in the form  
of a continuous, perforated or helical sleeve.
28. A polymeric product as in claim 27, characterized  
in that the thickness of the sleeve is from 0.005  
mm to 5.0 mm.
- 10 29. A polymeric product as in claims 27-28,  
characterized in that the sleeve is from 10 to 20  
mm in length.
30. A polymeric product as in claims 18-29,  
characterized in that the product is microporous.
- 15 31. A polymeric product as in claims 18-29,  
characterized in that the product is macroporous.
32. A polymeric product as in claims 18-31,  
characterized in that the polymeric material is  
disposed discontinuously on a mesh substrate.
- 20 33. A polymeric product as in claims 18-32,  
characterized in that the polymeric material  
contains living cells to facilitate healing.
- 25 34. A process for paving or sealing the interior  
surface of a tissue lumen comprising entering the  
interior of the tissue lumen and applying a layer  
of a polymer material to the interior surface of  
the tissue lumen, wherein the polymer is in  
intimate and conforming contact with the tissue  
surface.

35. A process as in claim 34, wherein the polymer is a biodegradable polymer.
36. A process as in claims 34-35, wherein the polymer is applied in a preshaped form to the interior surface of the vessel or organ.
37. A process as in claims 34-36, wherein the polymeric material contains a therapeutic agent.
38. The process as in claims 34-37, wherein multiple polymers containing various therapeutic agents are applied.
39. A process as in claims 34-38, characterized in that a catheter is used to position the polymeric material within the body.
40. A process according to claims 34-40, characterized in that a catheter in accordance with claims 1-17 is used to position the polymeric material within the body.
41. A process as in claims 34-41, characterized in that the polymeric material contains additives which promote organ regeneration.

## AMENDED CLAIMS

[received by the International Bureau on 15 January 1990 (15.01.90)  
original claims 4 and 6 cancelled; claims 1-3,5,7-11,13-17, 34,36 and  
40 amended; other claims unchanged (6 pages)]

1. A catheter device for application of a polymeric coating to a tissue lumen comprising a flexible tubular body having proximal and distal ends, which tubular body defines a lumen divided into a plurality of sublumens, each sublumen extending from the proximal end of the tubular body toward the distal end of the tubular body and connecting to at least one aperture in the tubular body whereby each sublumen forms a conduit for fluid flow between at least one aperture in the tubular body and the proximal end of the tubular body, characterized in that at least one of said sublumens is connected to a reservoir containing a monomer or prepolymer-containing liquid, whereby the polymeric coating is formed by delivering the monomer or prepolymer-containing liquid through the sublumen to the tissue lumen.
2. A catheter device as in claim 1, characterized in that the catheter includes at least one occluding balloon member disposed around the tubular body in alignment with the aperture of a second sublumen, whereby fluid flow through the second sublumen inflates the occluding balloon member.
3. A catheter device according to claims 1-2, characterized in that the device includes two occluding balloons, one disposed toward the proximal end of the tubular body from the aperture of the sublumen which is connected to the reservoir, and the other disposed toward the distal end of the tubular body from the aperture of the sublumen which controls delivery of polymeric coating material, whereby the occluding balloons together act to at least partially occlude a portion of the tissue lumen.

5. A catheter device for application of a polymeric coating to a tissue lumen comprising a flexible tubular body having proximal and distal ends, which tubular body defines a lumen divided into a plurality of sublumens, each sublumen extending from the proximal end of the tubular body toward the distal end of the tubular body and connecting to at least one aperture in the tubular body whereby each sublumen forms a conduit for fluid flow between at least one aperture in the tubular body and the proximal end of the tubular body, characterized in that the device includes an expansile member for delivery of at least partially preformed polymeric coating material to the tissue lumen and a heating element, said expansile member being disposed around the tubular body in alignment with an aperture of a first sublumen whereby the expansile member, upon expansion via fluid flow through the first sublumen, delivers polymeric coating material disposed on the surface of the expansile member to the tissue lumen where it is reconfigured by heat from the heating element to conform to the surface of the tissue lumen.
7. A catheter device as in claim 5, characterized in that a continuous, perforated or helical polymeric sleeve is disposed around the expansile member, whereby expansion of the balloon causes the polymeric sleeve to expand to coat the tissue lumen.
8. A catheter device as in claim 5, characterized in that the polymeric coating material is discontinuous and disposed on a mesh support.
9. A catheter device as in claims 1-3, characterized in that there is an expansile member disposed



around the tubular body in alignment with a further of said apertures and between the positioning balloons, whereby the thickness and surface configuration of the polymer coating is controlled by inflation of the expansile member.

10. A catheter device as in claim 9, characterized in that the expansile member has a contoured surface whereby a molded shape is imparted to the polymeric coating.
11. A catheter device as in claims 1-3 and 9-10, characterized in that the distal aperture of the sublumen controlling polymer delivery is shaped to provide flow acceleration.
12. A catheter device as in claim 11, characterized in that the shaped aperture includes a flow propellant device.
13. A catheter device as in claims 3 and 9-12, characterized in that inflation of the two occluding balloons is independently controllable.
14. A catheter device as in claims 1-3 and 9-13, characterized in that the device includes a heating element, whereby the polymeric coating material can be heated prior or subsequent to delivery to the tissue lumen.
15. A catheter device as in claims 1-3, 5 and 7-14, characterized in that the device includes an optical fiber whereby light can be delivered to the polymeric coating material.
16. A catheter device as in claims 1-3, 5, 7, 8 and 13-15 characterized in that the device includes a mechanically retractable protective sheath.

17. A catheter device as in claims 1-3, 5 and 7-17, characterized in that the polymeric material includes one or more therapeutic agents.
- 5 18. An at least partially preformed polymeric product for use in endoluminal paving and sealing.
19. A polymeric product as in claim 18, characterized in that the polymer employed is partially polymerized.
- 10 20. A polymeric product as in claim 19, characterized in that further polymerization of the polymeric product is thermally activated.
21. A polymeric product as in claim 19, characterized in that further polymerization of the polymeric product is photoactivated.
- 15 22. A polymeric product as in claims 18-21, characterized in that the polymer is biodegradable.
- 20 23. A polymeric product as in claim 22, characterized in that the product comprises polygluconolactone or a polyanhydride.
24. A polymeric product as in claims 18-21, characterized in that the product comprises polyethylene, polyvinyl chloride or ethylene vinyl acetate copolymer.
- 25 25. A polymeric product as in claims 18-24, characterized in that the polymer is mechanically deformable at temperatures achievable in vivo.

26. A polymeric product as in claims 18-25,  
characterized in that the product comprises one  
or more therapeutic or cell growth agents.
- 5 27. A polymeric product as in claims 18-26,  
characterized in that the product is in the form  
of a continuous, perforated or helical sleeve.
28. A polymeric product as in claim 27, characterized  
in that the thickness of the sleeve is from 0.005  
mm to 5.0 mm.
- 10 29. A polymeric product as in claims 27-28,  
characterized in that the sleeve is from 10 to 20  
mm in length.
30. A polymeric product as in claims 18-29,  
characterized in that the product is microporous.
- 15 31. A polymeric product as in claims 18-29,  
characterized in that the product is macroporous.
32. A polymeric product as in claims 18-31,  
characterized in that the polymeric material is  
disposed discontinuously on a mesh substrate.
- 20 33. A polymeric product as in claims 18-32,  
characterized in that the polymeric material  
contains living cells to facilitate healing.
- 25 34. A process for paving or sealing the interior  
surface of a tissue lumen characterized in that a  
polymeric material is introduced into the  
interior of the tissue lumen and reconfigured to  
form a layer of a polymer material on the  
interior surface of the tissue lumen, whereby the  
polymer is in intimate and conforming contact  
30 with the tissue surface.

35. A process as in claim 34, wherein the polymer is a biodegradable polymer.
36. A process as in claims 34-35, wherein the polymer is applied in a preshaped form to the interior  
5 surface of a vessel or organ.
37. A process as in claims 34-36, wherein the polymeric material contains a therapeutic agent.
38. The process as in claims 34-37, wherein multiple polymers containing various therapeutic agents  
10 are applied.
39. A process as in claims 34-38, characterized in that a catheter is used to position the polymeric material within the body.
40. A process according to claims 34-40,  
15 characterized in that a catheter in accordance with claims 1-3, 5 or 7-17 is used to position the polymeric material within the body.
41. A process as in claims 34-41, characterized in that the polymeric material contains additives  
20 which promote organ regeneration.

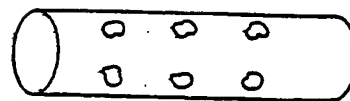


FIG. 1

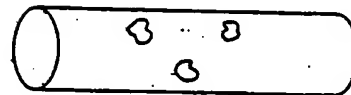


FIG. 2

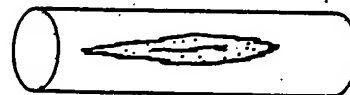


FIG. 3



FIG. 4

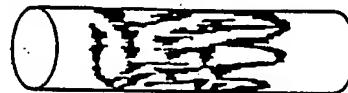
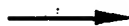
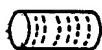


FIG. 5

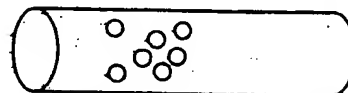


FIG. 6

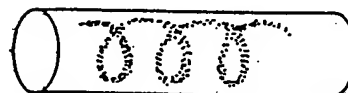


FIG. 7

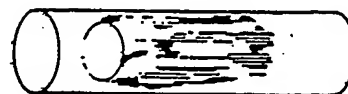


FIG. 8

SUBSTITUTE SHEET

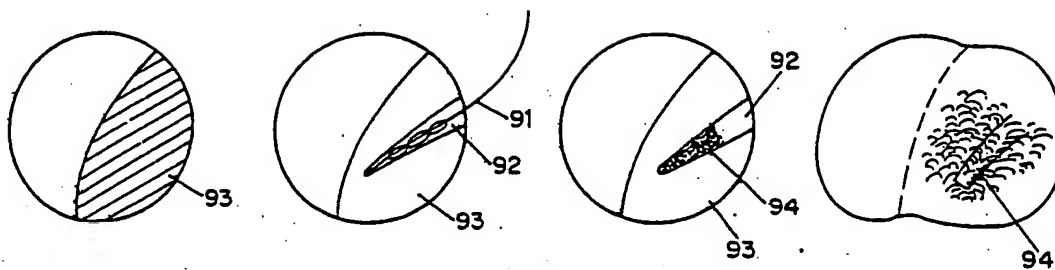


FIG. 9

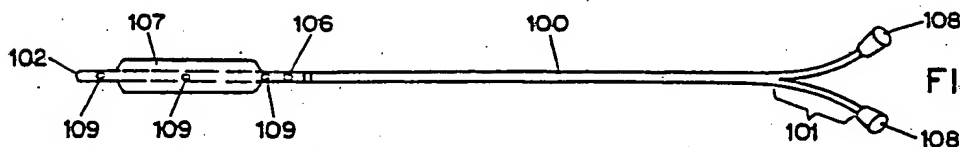


FIG. 10a

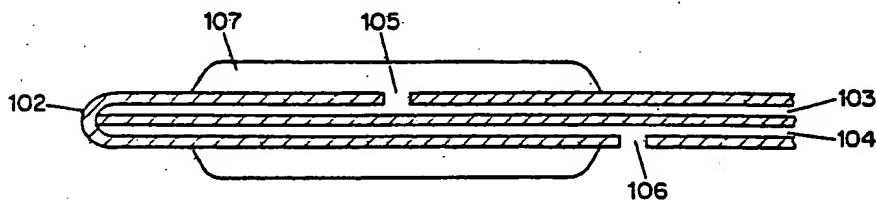


FIG. 10b

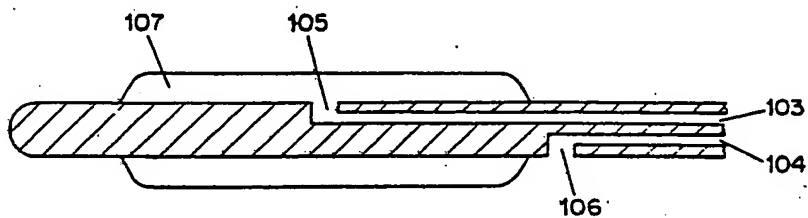


FIG. 10c

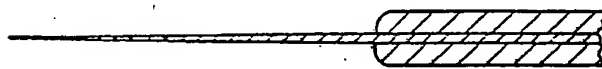


FIG. 10d

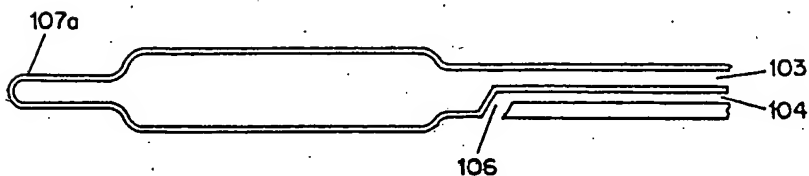


FIG. 10e

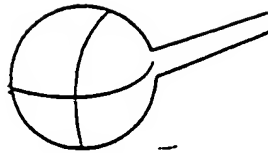


FIG. 11a

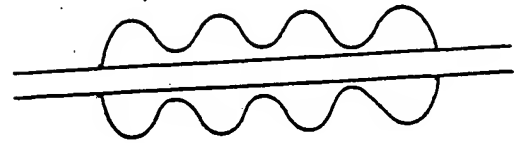


FIG. 11c

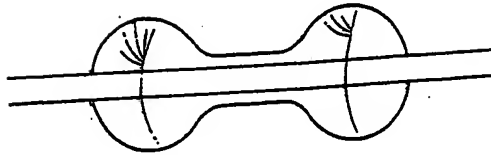


FIG. 11b

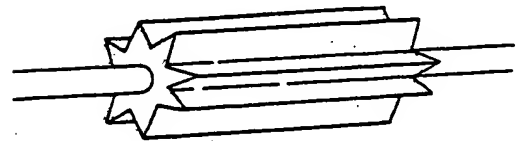


FIG. 11d

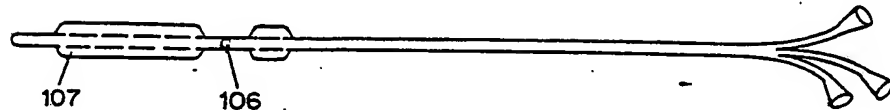


FIG. 12a

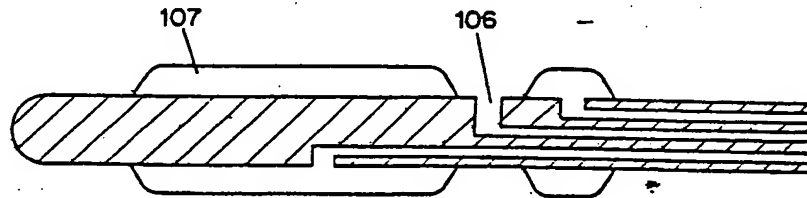


FIG. 12b

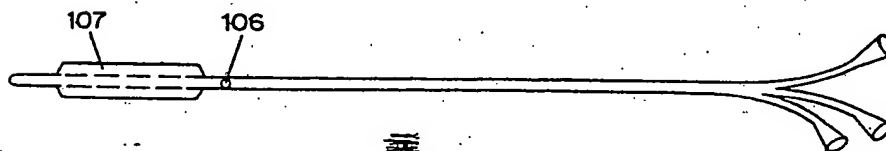


FIG. 12c

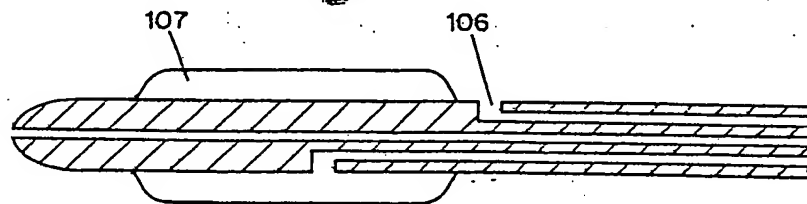


FIG. 12d

SUBSTITUTE SHEET

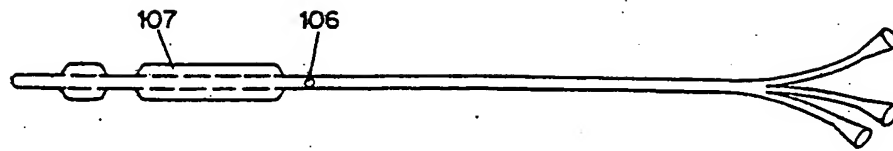


FIG. 12e

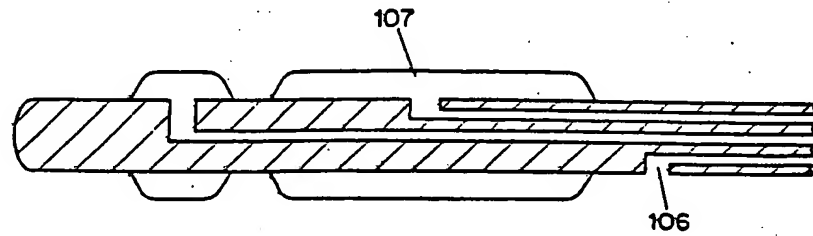


FIG. 12f

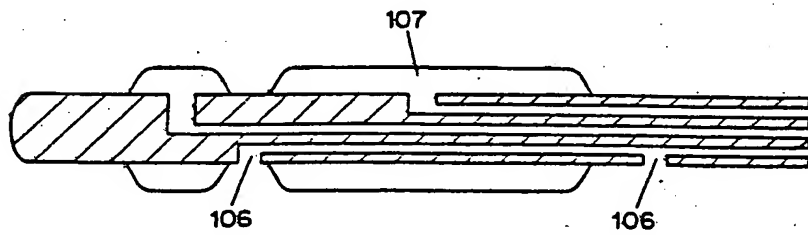


FIG. 12g

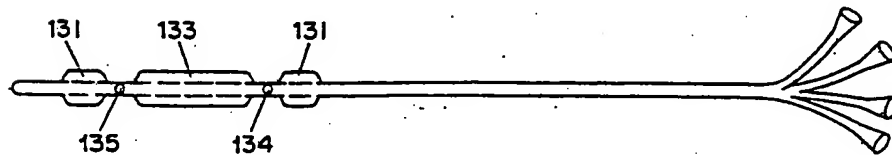


FIG. 13a

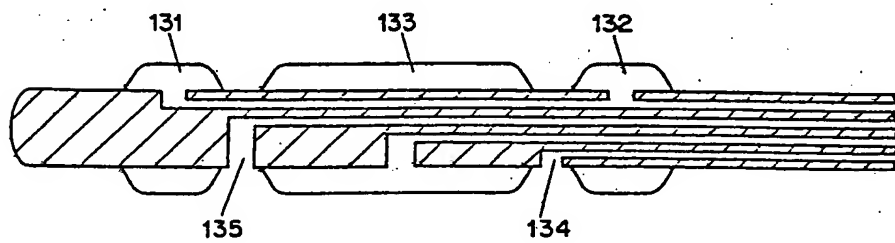


FIG. 13b

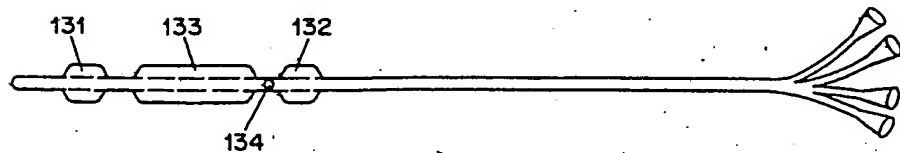


FIG. 13c

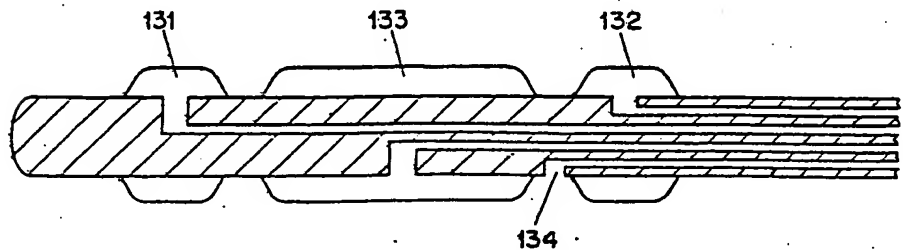


FIG. 13d



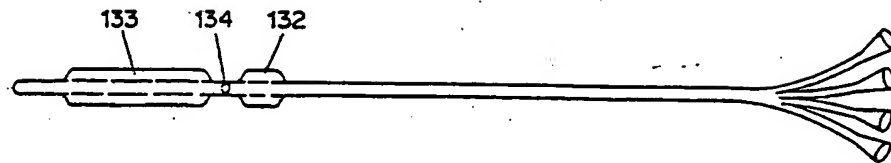


FIG. 13e

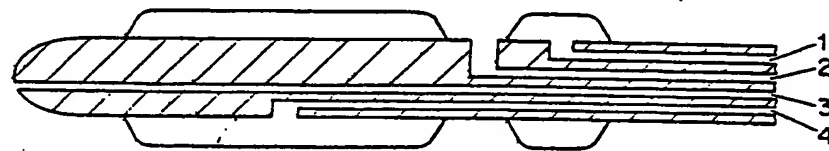


FIG. 13f

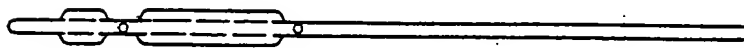


FIG. 13g

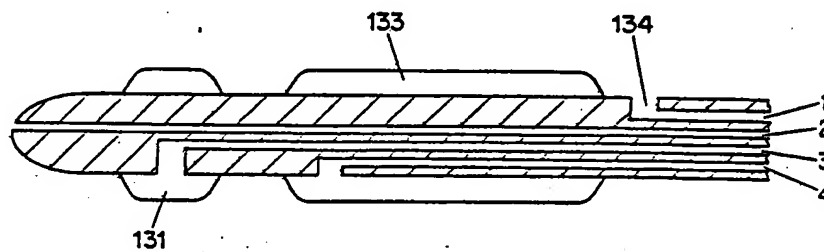


FIG. 13h

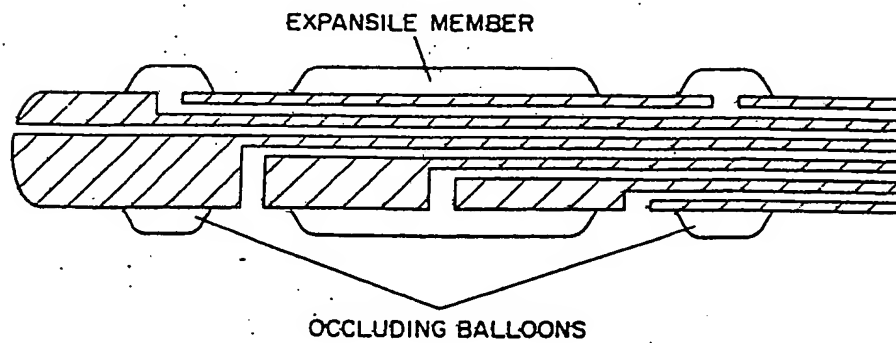
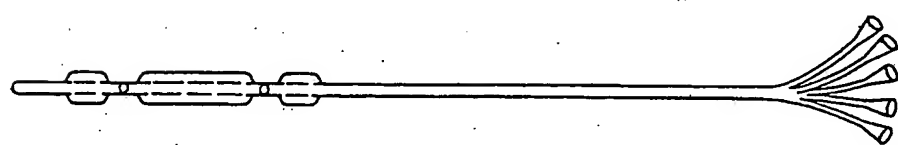


FIG. 14



FIG. 15a

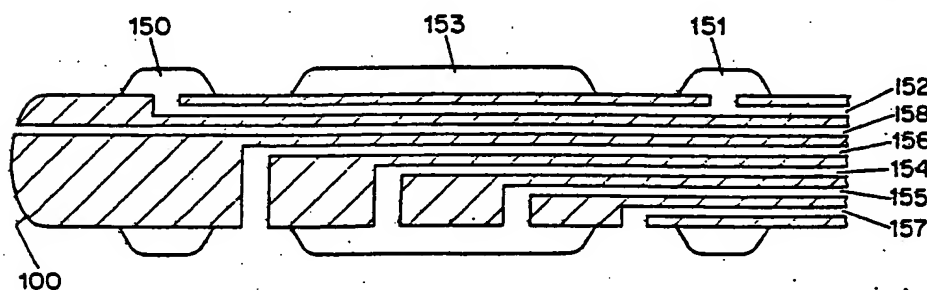


FIG. 15b

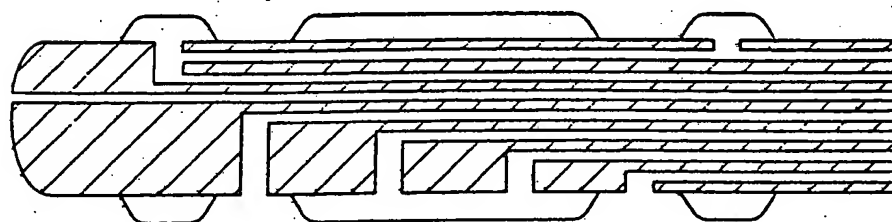
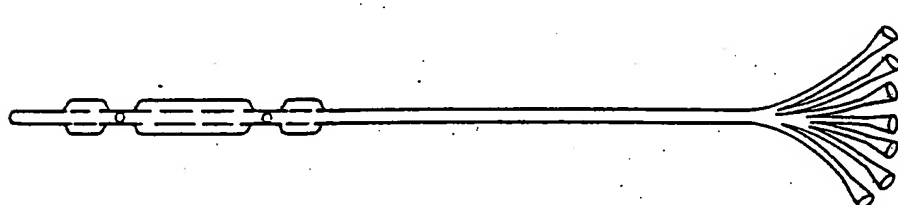


FIG. 16

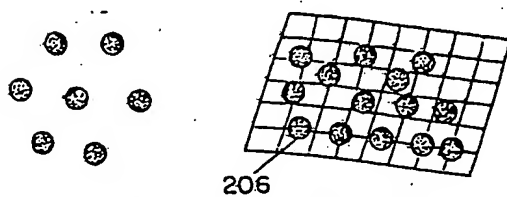


FIG. 20a

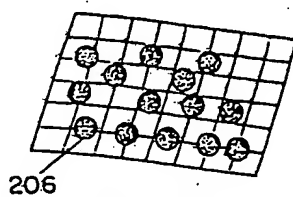


FIG. 20b

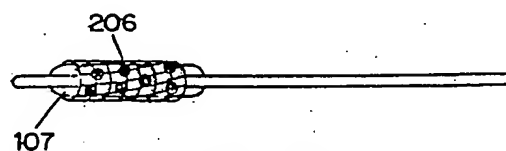


FIG. 20c

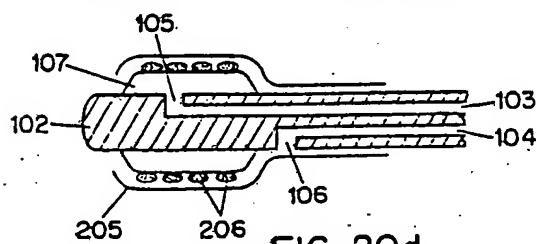


FIG. 20d

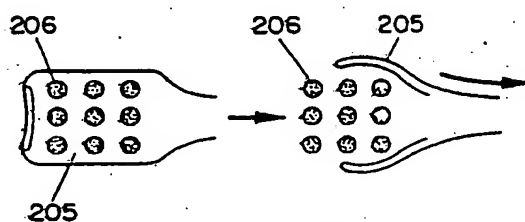


FIG. 20e

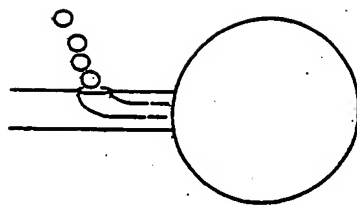


FIG. 21a

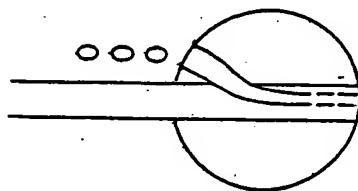


FIG. 21b

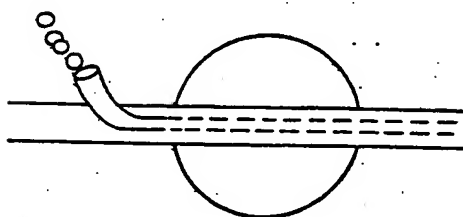


FIG. 21c

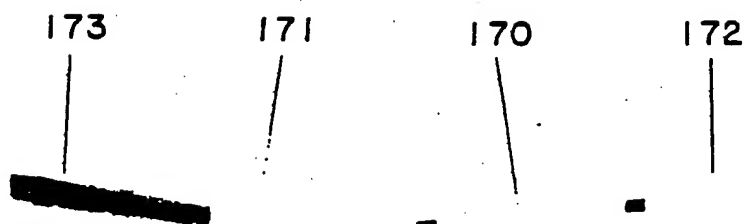


FIG. 17

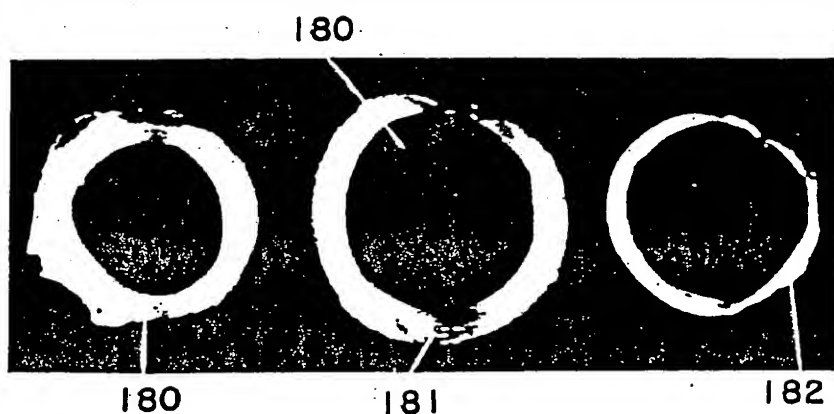


FIG. 18

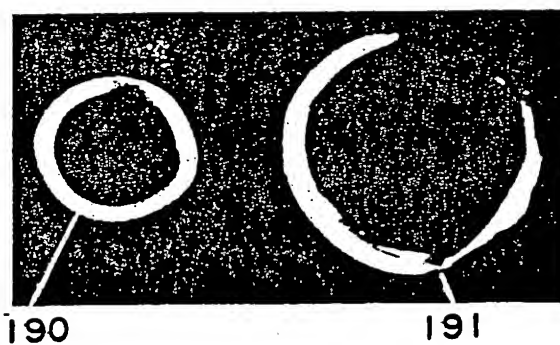


FIG. 19

SUBSTITUTE SHEET

# INTERNATIONAL SEARCH REPORT

International Application No. PCT/US89/03593

| <b>I. CLASSIFICATION OF SUBJECT MATTER</b> (If several classification symbols apply, indicate all) <sup>6</sup><br>According to International Patent Classification (IPC) or to both National Classification and IPC<br><b>IPC (4): A61M 29/00</b><br><b>U.S. Cl. 600/36; 623/1; 604/96, 101, 102</b>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         |                                                                                                                                                    |                                     |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------|-------------------------------------|------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------|------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------|-----|---|------------------------------------------------------------|---------|---|----------------------------------------------------------------------------------------|---------|---|--------------------------------------------------------------------------------------|----------|---|---------------------------------------------------------------------------------------|-------|---|-----------------------------------------------------------------------------|---------------|---|------------------------------------------------------------------------------------------------------|--------|
| <b>II. FIELDS SEARCHED</b><br><div style="text-align: center; border-top: 1px solid black; border-bottom: 1px solid black;">Minimum Documentation Searched <sup>7</sup></div> <table style="width: 100%; border-collapse: collapse;"> <tr> <th style="width: 20%; text-align: left; border-bottom: 1px solid black;">Classification System</th> <th style="text-align: left; border-bottom: 1px solid black;">Classification Symbols</th> </tr> <tr> <td style="vertical-align: top; border-right: 1px solid black; padding: 5px;">U.S.</td> <td style="padding: 5px;">604/96, 101, 102, 128; 623/1, 12, 66, 13; 600/36; 128/343, 344, 344R, 334C, 303R; 424/422, 423</td> </tr> </table> <div style="text-align: center; border-top: 1px solid black; border-bottom: 1px solid black;">Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched <sup>8</sup></div>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      |                                                                                                                                                    |                                     | Classification System                                                                    | Classification Symbols                                                                                                                             | U.S.                                                   | 604/96, 101, 102, 128; 623/1, 12, 66, 13; 600/36; 128/343, 344, 344R, 334C, 303R; 424/422, 423 |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| Classification System                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         | Classification Symbols                                                                                                                             |                                     |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| U.S.                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          | 604/96, 101, 102, 128; 623/1, 12, 66, 13; 600/36; 128/343, 344, 344R, 334C, 303R; 424/422, 423                                                     |                                     |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| <b>III. DOCUMENTS CONSIDERED TO BE RELEVANT <sup>9</sup></b><br><table style="width: 100%; border-collapse: collapse;"> <tr> <th style="width: 10%; text-align: left; border-bottom: 1px solid black;">Category <sup>9</sup></th> <th style="text-align: left; border-bottom: 1px solid black;">Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages <sup>12</sup></th> <th style="text-align: left; border-bottom: 1px solid black;">Relevant to Claim No. <sup>13</sup></th> </tr> <tr> <td style="vertical-align: top; border-right: 1px solid black; padding: 5px;">Y</td> <td style="padding: 5px;">US, A, 2,642,874 (KEELING) 04 June 1951<br/>See column 2, lines 8-32 and Figure 1.</td> <td style="vertical-align: top; padding: 5px;">1-4</td> </tr> <tr> <td style="vertical-align: top; border-right: 1px solid black; padding: 5px;">Y</td> <td style="padding: 5px;">US, A, 2,854,982 (PAGANO) 07 October 1958<br/>See Figure 2.</td> <td style="vertical-align: top; padding: 5px;">1-4, 13</td> </tr> <tr> <td style="vertical-align: top; border-right: 1px solid black; padding: 5px;">Y</td> <td style="padding: 5px;">US, A, 4,636,194 (WOLINSKY) 13 January 1987<br/>See column 2, lines 32-68 and Figure 1.</td> <td style="vertical-align: top; padding: 5px;">1-4, 13</td> </tr> <tr> <td style="vertical-align: top; border-right: 1px solid black; padding: 5px;">Y</td> <td style="padding: 5px;">US, A, 4,739,762 (PALMAZ) 26 April 1988<br/>See column 1, line 60-column 14, line 56.</td> <td style="vertical-align: top; padding: 5px;">5-10, 16</td> </tr> <tr> <td style="vertical-align: top; border-right: 1px solid black; padding: 5px;">X</td> <td style="padding: 5px;">US, A, 4,674,506 (ALCOND) 23 June 1987<br/>See Figures 1, 7 and column 4, lines 36-42.</td> <td style="vertical-align: top; padding: 5px;">34-36</td> </tr> <tr> <td style="vertical-align: top; border-right: 1px solid black; padding: 5px;">X</td> <td style="padding: 5px;">US, A, 4,140,126 (CHOUDHURY) 20 February 1979<br/>See column 4, lines 27-53.</td> <td style="vertical-align: top; padding: 5px;">34, 36 and 39</td> </tr> <tr> <td style="vertical-align: top; border-right: 1px solid black; padding: 5px;">Y</td> <td style="padding: 5px;">US, A, 4,702,917 (SCHINDLER) 27 October 1987<br/>See column 7, lines 11-41 and column 8, lines 31-35.</td> <td style="vertical-align: top; padding: 5px;">37, 38</td> </tr> </table> |                                                                                                                                                    |                                     | Category <sup>9</sup>                                                                    | Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages <sup>12</sup>                                     | Relevant to Claim No. <sup>13</sup>                    | Y                                                                                              | US, A, 2,642,874 (KEELING) 04 June 1951<br>See column 2, lines 8-32 and Figure 1. | 1-4 | Y | US, A, 2,854,982 (PAGANO) 07 October 1958<br>See Figure 2. | 1-4, 13 | Y | US, A, 4,636,194 (WOLINSKY) 13 January 1987<br>See column 2, lines 32-68 and Figure 1. | 1-4, 13 | Y | US, A, 4,739,762 (PALMAZ) 26 April 1988<br>See column 1, line 60-column 14, line 56. | 5-10, 16 | X | US, A, 4,674,506 (ALCOND) 23 June 1987<br>See Figures 1, 7 and column 4, lines 36-42. | 34-36 | X | US, A, 4,140,126 (CHOUDHURY) 20 February 1979<br>See column 4, lines 27-53. | 34, 36 and 39 | Y | US, A, 4,702,917 (SCHINDLER) 27 October 1987<br>See column 7, lines 11-41 and column 8, lines 31-35. | 37, 38 |
| Category <sup>9</sup>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         | Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages <sup>12</sup>                                     | Relevant to Claim No. <sup>13</sup> |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| Y                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             | US, A, 2,642,874 (KEELING) 04 June 1951<br>See column 2, lines 8-32 and Figure 1.                                                                  | 1-4                                 |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| Y                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             | US, A, 2,854,982 (PAGANO) 07 October 1958<br>See Figure 2.                                                                                         | 1-4, 13                             |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| Y                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             | US, A, 4,636,194 (WOLINSKY) 13 January 1987<br>See column 2, lines 32-68 and Figure 1.                                                             | 1-4, 13                             |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| Y                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             | US, A, 4,739,762 (PALMAZ) 26 April 1988<br>See column 1, line 60-column 14, line 56.                                                               | 5-10, 16                            |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| X                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             | US, A, 4,674,506 (ALCOND) 23 June 1987<br>See Figures 1, 7 and column 4, lines 36-42.                                                              | 34-36                               |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| X                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             | US, A, 4,140,126 (CHOUDHURY) 20 February 1979<br>See column 4, lines 27-53.                                                                        | 34, 36 and 39                       |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| Y                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             | US, A, 4,702,917 (SCHINDLER) 27 October 1987<br>See column 7, lines 11-41 and column 8, lines 31-35.                                               | 37, 38                              |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| <div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p><sup>10</sup> Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"A" document member of the same patent family</p> </div> </div>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            |                                                                                                                                                    |                                     |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| <b>IV. CERTIFICATION</b><br><table style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%; border-right: 1px solid black; padding: 5px;">           Date of the Actual Completion of the International Search<br/><br/> <b>06 November 1989</b> </td> <td style="width: 50%; padding: 5px;">           Date of Mailing of this International Search Report<br/><br/> <div style="text-align: center; font-size: 1.2em; font-weight: bold;">21 NOV 1989</div> </td> </tr> <tr> <td style="width: 50%; border-right: 1px solid black; padding: 5px;">           International Searching Authority<br/><br/> <b>ISA/US</b> </td> <td style="width: 50%; padding: 5px;">           Signature of Authorized Officer<br/> <b>A. Cannon - Primary Examiner</b> </td> </tr> </table>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             |                                                                                                                                                    |                                     | Date of the Actual Completion of the International Search<br><br><b>06 November 1989</b> | Date of Mailing of this International Search Report<br><br><div style="text-align: center; font-size: 1.2em; font-weight: bold;">21 NOV 1989</div> | International Searching Authority<br><br><b>ISA/US</b> | Signature of Authorized Officer<br><b>A. Cannon - Primary Examiner</b>                         |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| Date of the Actual Completion of the International Search<br><br><b>06 November 1989</b>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      | Date of Mailing of this International Search Report<br><br><div style="text-align: center; font-size: 1.2em; font-weight: bold;">21 NOV 1989</div> |                                     |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |
| International Searching Authority<br><br><b>ISA/US</b>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        | Signature of Authorized Officer<br><b>A. Cannon - Primary Examiner</b>                                                                             |                                     |                                                                                          |                                                                                                                                                    |                                                        |                                                                                                |                                                                                   |     |   |                                                            |         |   |                                                                                        |         |   |                                                                                      |          |   |                                                                                       |       |   |                                                                             |               |   |                                                                                                      |        |

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